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(54) **Water sensitive hot melt adhesives for nonwoven applications**

Heisschmelze wasserempfindliche Klebstoffe für Verwendungen eines nichtgewebten Stoffes

Thermoadhésifs sensibles à l'eau pour des applications en étoffe non-tissée

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Description

A nonwoven fabric is defined as an interlocking polymer network characterized by flexibility, porosity and integrity. The individual fibers used to compose the nonwoven fabric may be synthetic, naturally occurring, or a combination of the two. The individual fibers may be mechanically, chemically, or thermally bonded to each other. Nonwovens are used commercially for a variety of applications including insulation, packaging, household wipes, surgical drapes, medical dressings, and in disposable articles such as diapers, adult incontinent products and sanitary napkins.

In many of the aforementioned applications it is necessary to adhere the nonwoven to another substrate or component. The second substrate may be another nonwoven, or an unrelated material. A commonly employed technique to bond the assembly together is the use of a hot melt adhesive. Hot melt adhesives allow for cost and time efficient manufacturing since there is no evaporation step necessary as is the case for water based or solvent based adhesive systems. Suitable hot melt adhesives must possess excellent adhesion to the substrates involved. For nonwoven applications they must also possess good flexibility (or hand), no staining or bleed through, suitable viscosity, set speed and open time to function on commercially available equipment and finally, acceptable thermal aging properties.

Recently a variety of nonwoven applications have been developed which require that the hot melt adhesive demonstrate appreciable water solubility, dispersibility or sensitivity. In these situations the hot melt adhesive must provide a durable bond to the nonwoven until exposed to a predetermined condition (e.g., room temperature water), after which the adhesive would release from the substrate(s). This ambient water releasability is a particularly desirable property in the disposable market where flushability and/or degradeability are becoming critical.

It has now been discovered that a specific class of hot melt adhesives comprising 50-100 weight % of a graft copolymer comprising 40-85 weight percent of at least one vinyl monomer and 15-60 weight percent of at least one polyalkylene oxide polymer and 0-50 weight % of a compatible tackifying resin, is desirable for bonding nonwoven fabric substrates to themselves or to other substrates, yet will later release upon exposure to ambient temperature water. These adhesives are characterized by a glass-transition temperature (T_g) of -40°C to $+40^{\circ}\text{C}$ as determined by the maximum value of G''/G' or $\tan \delta$ when measured using the Rheometrics Dynamic Spectrometer at 10 radians per second.

Accordingly, the invention relates to a process for bonding nonwoven fabric substrates to similar or dissimilar substrates comprising the steps of applying to at least one substrate a molten hot melt adhesive composition, said adhesive having a T_g of -40 to $+40^{\circ}\text{C}$ comprising 50-100 weight % of a graft copolymer comprising (a) 40-85 weight percent of at least one vinyl monomer and (b) 15-60 weight percent of at least one polyalkylene oxide polymer having a molecular weight greater than 3000 and a polymerized ethylene oxide content of at least 50% by weight, and 0-50 weight % of a compatible tackifying resin; and bonding another substrate thereto.

In preferred embodiments, at least one substrate is a nonwoven and the adhesive comprises 50-95 weight % of the graft copolymer and 5-50 weight % of a compatible tackifying resin.

The graft copolymers used herein are actually mixtures of graft and homopolymers of the respective monomers. For convenience these graft copolymer/homopolymer mixtures will be referred to as "graft copolymers" throughout the specification and claims. The preferred vinyl monomer used in the graft copolymer is vinyl acetate or an alkyl-substituted acrylate such as methyl acrylate or ethyl acrylate or mixtures thereof. The preferred polyalkylene oxide is a polyethylene oxide homo- or copolymer with a molecular weight of 3,000-100,000. Preferred tackifying resins for those adhesives which are tackified include rosin, hydrogenated rosin, rosin esters, terpene phenolic and dimerized rosins. As noted above, the use of tackifying resin is optional and may in fact prove undesirable for a number of applications. Other conventional additives such as plasticizers, antioxidants, pigments, dyes and waxes may be optionally incorporated into the adhesives.

The specific formulation of the water sensitive hot melt adhesives will vary depending upon the intended end use thereof. Thus, adhesives comprising 100% of the graft copolymer are particularly useful for bonding superabsorbent particulate materials onto nonwoven substrates where releasability of the superabsorbent is desired. These untackified hot melt adhesives are also highly suitable for creping adhesives.

On the other hand, highly tackified graft copolymer hot melt adhesives, particularly those prepared from the higher molecular weight (e.g., those having a viscosity greater than about 5000 mPas (cps) at 175°C) graft copolymers are especially suited for end uses where toughness and adhesion as well as flexibility are desired. As such, they find use as construction adhesives for disposable diapers where water sensitivity at relatively low temperatures is desirable so as to provide release from the various substrates to facilitate flushability of all or of a portion of the diaper.

Other hot melt adhesive compositions containing intermediate levels of tackifier are also useful as adhesives in other areas of nonwoven manufacture including in less demanding construction adhesives, nonwoven garment construction, disposable wipes and the like.

The specific water-soluble polyalkylene oxide polymers used in the graft copolymers described herein have a molecular weight of 3,000 to 100,000 and a polymerized ethylene oxide content of at least 50% by weight. The polyalkylene oxide polymers may be homopolymers of ethylene oxide (including the ester and ether derivatives thereof),

random copolymers of ethylene and propylene oxide, block copolymers of ethylene and propylene oxides, or mixtures thereof. It will be noted that mixtures of different polyalkylene oxide polymers may be utilized, and copolymers and homopolymers may be used together in such mixtures. The polymers are commercially available from companies such as Union Carbide (the polyethylene oxide polymers, poly (ethylene oxide/propylene oxide) copolymers and monomethyl
 5 ethers of polyethylene oxide), BASF Wyandotte (the block copolymers) and Dow Chemical Company (the homopolymers and random copolymers). More specifically, the Union Carbide products sold under the trade names CARBOWAX (for polyethylene oxides) and METHOXY CARBOWAX (for the monomethyl ethers of polyethylene oxide) have an average molecular weight roughly indicated by the numeral following the trade name.

The polymerized ethylene oxide content of the polyalkylene oxide polymer should be at least 50% by weight of the polymer and preferably at least 75%. Polymers having a lower content of polymerized ethylene oxide groups display only limited solubility in water, and hence are not useful as the water-sensitive polymer component of the present invention. Polyalkylene oxide polymers containing less than about 50% by weight of polymerized ethylene oxide are, however, useful as plasticizers or diluents.

The polyalkylene oxide polymers having a molecular weight of less than about 3,000, when used as the sole
 15 polyalkylene oxide, have been found not to impart water dispersibility to the graft copolymers, and hence, such lower molecular weight polyalkylene oxide fractions should not be included in determining the proportion of the water-soluble component present in the graft copolymer. On the other hand, such low molecular weight polyalkylene oxides (e.g., CARBOWAX 600) are useful as plasticizers or diluents.

The vinyl monomers useful in the graft copolymers of the present invention are preferably vinyl acetate and the
 20 lower alkyl-substituted acrylates or methacrylates such as methyl acrylate and ethyl acrylate. Other useful vinyl monomers include the alkyl esters of acrylic acid containing 1 to 10 carbon atoms in the alkyl portion; styrene; and vinyl esters such as vinyl propionate, vinyl butyrate, vinyl 2-ethylhexanoate and the like. Use of the vinyl monomers, and in particular the vinyl acetate monomer, provides sufficient chain transfer in grafting to produce a graft copolymer which is thermally stable, moderately polar and readily formulatable into an adhesive. The graft copolymer preferably comprises from 40 to 85% of at least one vinyl monomer and about 15-60% by weight of at least one water-soluble poly-
 25 alkylene oxide polymer, and most preferably contains 15 to 45% by weight of the polyalkylene oxide component(s).

While various vinyl monomers may be utilized by themselves for grafting onto the water-soluble polymer backbone, small amount of other ethylenically unsaturated monomers may be utilized as comonomers with the vinyl monomer to improve particular properties such as water-dispersibility, adhesion, softness and the like. Monomers useful as comonomers with the vinyl monomers include 2-hydroxyethyl acrylate, N-vinyl pyrrolidone, sodium vinyl sulfonate (the sodium
 30 salt of ethylene sulfonic acid) and the alkyl esters of methacrylic acid containing 1-8 carbon atoms in the alkyl portion. Such comonomers are generally utilized in quantities not exceeding about 40% by weight of the total graft copolymer.

The graft copolymers used herein as well as a process for the production thereof is described in US-A-3,891,584, the disclosure of which is incorporated by reference.

In addition to the water-soluble polyalkylene oxide polymer or polymers, vinyl monomer or monomers and optional
 35 ethylenically unsaturated monomers, the adhesive may also contain one or more compatible tackifying resins in amounts up to about 50% by weight. Such resins primarily produce a reinforcing effect or plasticizing (flexibility) effect, but also contribute stickiness, particular wetting ability, and viscosity control to the graft copolymer. Exemplary of such tackifying resins are rosin (from gum, wood or tall oil) and the rosin derivatives, the phenolic modified coumarone indene resins (sold by Neville Chemical Company of Neville Island, Pennsylvania under the trade name of NEVILLAC),
 40 the coumarone indene resins with softening points of about 5° to 117°C (sold by the aforementioned Neville Chemical Company under the trade name CUMAR, the phenolic modified terpene resins (sold by Arizona Chemical Company, Inc. of Elizabeth, New Jersey under the NIREZ trade name). (The softening points of the resins referred to above are Ball and Ring softening points determined using method ASTM E28-58T.) As noted above and as illustrated in the
 45 Examples presented herein, particularly preferred adhesives for certain applications are prepared using levels of tackifier in excess of about 25% by weight.

The adhesive may also contain small amounts, i.e., up to about 25% by weight, of a compatible plasticizing oil and/or wax diluent. Particularly useful are diluents selected from the group consisting of glycerol mono- and distearate, synthetic long chain linear polymeric alcohols, stearic acid, high acid number waxlike materials of mono- or dicarboxylic
 50 acids, acid wax derived from montan wax, stearyl alcohol, hydrogenated castor oil, ethoxylated alcohols, 12(OH) stearic acid and stearic-cetyl alcohol.

The resulting adhesives of the present invention are characterized by their ability to provide a durable bond to a nonwoven article and otherwise meet the unique requirements of the application (including flexibility, non-staining, and machinable viscosity) and later release upon exposure to water at ambient temperatures after a desired residence
 55 period. The adhesives described herein also possess exceptional thermal stability which distinguishes them from other moisture sensitive technologies. Additionally, many of the compositions described herein display long open time and pressure sensitivity necessary for spray or melt blown application methods, yet later (e.g., after several hours or days), slowly crystallize. This crystallization results in a loss of surface tack but does not detract appreciably from adhesion.

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In addition, the crystallization actually enhances properties such as elevated temperature peel and shear resistance and resistance to strike-through.

The adhesive product can be applied to a substrate such as a nonwoven article by a variety of methods including coating or spraying in an amount sufficient to cause the article to adhere to another substrate such as nonwoven, or an unrelated material such as a low density polyolefin or other conventionally employed water impermeable substrate. When exposed to water of a specific temperature for a specific period of time, the nonwoven releases.

The following examples illustrate the production of suitable hot melt adhesives as well as the use thereof in a variety of applications. In the examples, all parts are by weight and all temperatures in degree Celsius unless otherwise noted. Test procedures used herein are as follows:

Test Procedures

Melt viscosities of the hot melt adhesives/binders were determined on a Brookfield RVT Thermosel viscometer using a number 27 spindle at 20 or 50 rpms.

The heat stability of the adhesive/binder samples was determined in the following manner: a 60 gram sample was stored in a covered jar at 175°C for 72 hours and observed for the formation of gel, skin or any deleterious change in color. The viscosity change upon this thermal aging was then determined as stated above.

The glass transition temperature, T_g, of the samples was determined using a Rheometrics Dynamic Analyzer (RDA-700). Specifically, 0.15-0.20 gram of the sample was placed between 8 mm parallel plates at a separation of 2.0 - 2.5 mm. The sample was then cooled to approximately -80°C and then heated at 5°C/minute to >60°C while stressed in a sinusoidal manner at a rate of 10 radians/second. The strain was adjusted throughout the test from 0.03% to a maximum of 10%. The temperature corresponding to the maximum value of the ratio of loss modulus G'' to storage modulus G' (this ratio is also referred to as Tan Delta) is recorded as the glass transition temperature, T_g.

Polypropylene nonwoven to nonwoven bonds used in testing nonwoven adhesion and water sensitivity were prepared by applying a 1.0 - 1.5 mm wide (compressed) bead of molten hot melt at 177 - 191°C (350 - 375°F) to one piece of 5 cm x 15 cm (2" x 6") nonwoven and mating it to a second piece of nonwoven of the same size. Open time was <2 seconds and dwell time 10 seconds. These bonds were then aged at 21°C (70°F)/50% RH.

Nonwoven adhesion of the various samples was determined by pulling polypropylene nonwoven to polypropylene nonwoven bonds apart in a peel mode at 12"/min on an Instron tester. The bond is characterized by the presence or absence of fiber tear (F.T.) and/or the amount of resistance (N/cm (lbs/in)). If F.T. was noted on all three samples, the amount of resistance was not recorded. Conventional hot melt adhesives used for bonding nonwovens typically display fiber tearing bonds (F.T.) and/or resistance ≥ 0.35 N/cm (≥ 0.2 lb/in).

Water sensitivity of nonwoven bonds prepared with the hot melt adhesive/binder samples was determined by immersing polypropylene nonwoven to polypropylene nonwoven (which had been aged 1 week at 21°C (70°F)/50% RH) in water at various temperatures for various lengths of time. After a specified time interval (e.g., 15 minutes) a nonwoven to nonwoven bond was removed from the water and stressed by hand in a peel mode. The bond is characterized as fiber tearing (F.T.) or non-fiber tearing (no F.T.). Conventional hot melt adhesives used for nonwovens would produce fiber tearing bonds and/or bonds with good resistance both before and after immersion. The adhesives used in the present invention are characterized by various levels of water sensitivity which result in a reduction in bond strength to nonwoven after immersion in water for a specified time interval and/or at a specified temperature.

180° T Peel Testing Procedure

The samples are prepared as follows. A glue line was extruded onto nonwoven at approximately 149-163°C (300-325°F) with a line speed of 30 m/min (100 FPM) to form a glue line approximately 0.5 mm wide with a coating weight of approximately 1.6 - 2.4 mg/cm (4-6 mg/linear inch). A second nonwoven substrate was immediately bonded to the glue bead with bonding pressure of approximately 415 kPa (60 psi). Samples were then cut parallel to adhesive lines, leaving at least 0.3125 cm (1/8") on each side of the exterior adhesive lines. The samples were conditioned overnight at 21°C (70°F)/50% RH constant temperature and humidity.

Instron Testing: The ends of each sample were taped, then placed in jaws, with the adhesive coated nonwoven in the stationary jaw. The sample was pulled at 30 cm/min (12 in/min) crosshead speed, 5 cm/min (2 in/min) chart speed in 180° T peel mode and the average peel value recorded in grams or pounds for each product tested. The deformation and the type of failure (adhesive, cohesive) were also recorded.

Example

A useful graft copolymer of 30 parts poly (ethylene oxide) (PEG) and 70 parts vinyl acetate (VA) was prepared by charging the following ingredients into a one liter flask:

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Polyglycol E-8000 (Dow Chemical Co.)	150 g
t-butyl peroxyperbenzoate	0.4 g

The flask was fitted with a stainless steel stirrer, thermometer, condenser, nitrogen inlet tube and dropping funnel containing 350 g of vinyl acetate. 45 ml of the vinyl acetate was added and the mixture was heated to reflux. 1 ml of a 10% solution of 70% benzoyl peroxide in ethyl acetate was added to initiate the polymerization. The remainder of the vinyl acetate was added over a three hour period. The reaction mixture was heated to 155 to 160°C and held at this temperature for 15 minutes. 1.5 gram of Irganox 1010 antioxidant (Ciba-Geigy) was added and the residual monomer was removed by vacuum distillation. The viscosity of the product was 2,525 mPas (cps) at 177°C (350°F).

Similar graft copolymers were prepared using other proportions of vinyl acetate and PEG 8000 as shown in Table I. The resultant graft copolymers were then tested "neat" as adhesives for nonwovens. The results of the testing are also shown in Table I.

Table I

	1	2	3	4
60 vinyl acetate/40 PEG-8000 copolymer	100.0	----	----	----
70 vinyl acetate/30 PEG-8000 copolymer	----	100.0	----	----
80 vinyl acetate/20 PEG-8000 copolymer	----	----	100.0	----
90 vinyl acetate/10 PEG-8000 copolymer	----	---	---	100.0
Viscosity @ 177°C (350°F)	1310 mPas (cps)	2500 mPas (cps)	6475 mPas (cps)	13,775 mPas (cps)
Glass Transition Temp (Tg)	-8°C	+ 5°C	+ 20°C	+36°C
Nonwoven Adhesion	F.T.	F.T.	F.T.	F.T.
Water Sensitivity				
15 min/21°C (70°F)	No F.T.	No F.T.	F.T.	F.T.
24 hrs/21°C (70°F)	No F.T.	No F.T.	F.T.	F.T.
15 min/38°C (100°F)	No F.T.	No F.T.	F.T.	F.T.
24 hrs/38°C (100°F)	No F.T.	No F.T.	No F.T.	F.T.
Heat Stability (72hrs/177°C (350°F))				
Color	Amber	Dark Amber	Dark Amber	Dark Brown
Gel	None	None	None	None
Skin	None	None	None	None
% Viscosity Change	-38.9%	-7.5%	-7.3%	+3.2%

The results presented in Table 1 demonstrate the suitability of the untackified graft copolymers containing greater than 10% grafted polyalkylene oxide and characterized by glass transition values of between -40°C and +40°C for providing a fiber tearing bond to nonwoven, which bond also demonstrates water sensitivity either at 21°C or 38°C (70°F or 100°F). These unformulated copolymers also demonstrate good heat stability (no skin or gel is formed).

In the following examples, various graft copolymers were formulated with various tackifying resins and/or oils and tested as adhesives for non-woven substrates.

TABLE 2

	5	6	7	8	9	10	11	12
60 vinyl acetate/25 PEG-8000/15 Ucon 75H-90,000	100.0	70.0	---	---	---	---	---	---

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TABLE 2 (continued)

	5	6	7	8	9	10	11	12
5	Brazilian Gum Rosin	---	30.0	---	20.0	30.0	---	---
	35 vinyl acetate/ 35 vinyl 2-ethylhexanoate/ 25 Ucon 75H-90,000 15 PEG- 8000	---	---	100.0	80.0	70.0	75	---
10	Permalyn 85	---	---	---	---	25	---	---
	Dymere	---	---	---	---	---	25	---
	Stereon 840A	---	---	---	---	---	---	21
	Zonatac 105	---	---	---	---	---	---	60
15	Kaydol USP White Mineral Oil	---	---	---	---	---	---	19
20	Viscosity at 177°C (350°F)	3275 mPas (cps)	920 mPas (cps)	9500 mPas (cps)	2810 mPas (cps)	1855 mPas (cps)	3763 mPas (cps)	3800 mPas (cps)
	Glass Transition Temp. (T _g)	-5°C	+10°C	+4°C	+3°C	+8°C	+22°C	+11°C
25	Nonwoven Adhesion (pp/ pp-12"/min)	F.T.	F.T.	Not Tested	0.88 N/ cm (0.5 lb/in)	F.T.	0.88 N/ cm (0.5 lb/in)	1.4 N/cm (0.8 lb/ in) F.T.
30	LDPE/PP Nonwoven 180° T-peel (12"/min)	No F.T. 0.18 N/ cm (0.1 lb/ in)	No F.T. 0.53 N/ cm (0.3 lb/in)	No F.T. 0.18 N/cm (0.1 lb/ in)	F.T. (1 bond) 0.44 N/ cm (0.25 lb/in) (2 bonds)	No F.T. 0.88 N/ cm (0.5 lb/in)	F.T. (1 bond) 0.70 N/ cm (0.4 lb/in) (2 bonds)	F.T.
35	Water (pp/pp) Sensitivity			Not Tested				
	15 min/21°C (70°F)	No F.T.	F.T.		No F.T.	F.T.	No F.T.	No F.T.
	24 hr/21°C (70°F)	No F.T.	No F.T.		No F.T.	F.T.	No F.T.	No F.T.
40	15 min/38°C (100°F)	No F.T.	F.T.		No F.T.	F.T.	No F.T.	No F.T.
	24 hr/38°C (100°F)	No F.T.	No F.T.		No F.T.	F.T.	No F.T.	No F.T.

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TABLE 2 (continued)

	5	6	7	8	9	10	11	12
Heat Stability (72 hrs/ 177°C (350°F))			Not Tested					
Color	Black	Amber		Dark Brown	Black	Brown	Dark Brown	Light Yellow
Gel	None	None		None	None	None	None	None
Skin	None	None		None	None	None	None	None
% viscosity change	-65.3%	-29.9%		-66.9%	-59.6%	-50.0%	-67.1%	-24.8%
<p>Ucon 75H-90,000 is an ethylene oxide, propylene oxide copolymer containing approximately 75% ethylene oxide available from Union Carbide.</p> <p>Permalyne 85 is a glycerol ester of rosin available from Hercules Inc.</p> <p>Dymere is a polymerized rosin available from Hercules, Inc.</p> <p>Stereon 840A is a styrene butadiene block copolymer available from Firestone.</p> <p>Zonatac 105 is a styrenated polyterpene resin from Arizona Chemicals.</p> <p>Kaydol USP White mineral oil is available from Witco.</p>								

The results presented in Table 2 demonstrate the suitability of a graft terpolymer (5) based on vinyl acetate, polyethylene oxide and polypropylene oxide/ethylene oxide for producing useful water sensitive bonds to both the nonwoven/nonwoven and nonwoven/LDPE constructions as contrasted with a commercially available block copolymer based hot melt adhesive (12). Incorporation of a compatible resin (6) increases the adhesion to low density polyethylene and is also useful. Also demonstrated in Table 2 is the use of a graft copolymer (7) based on vinyl acetate, vinyl-2-ethylhexanoate, polyethylene oxide, and polypropylene oxide/ethylene oxide. Although the graft copolymer (7) or similar copolymer compositions may prove useful without further formulation, formulating with compatible resins appears to improve the utility of the composition. Some properties such as LDPE/nonwoven adhesion and viscosity are improved by the addition of compatible resins such as gum rosin at levels such as 20 and 30 percent (8 and 9). Also useful as compatible resins are rosin ester (10) and dimerized rosin (11).

Example 3

A graft copolymer of 70 weight percent vinyl acetate and 30 weight percent polyethylene oxide of 8000 molecular weight was prepared as described in Example 1.

TABLE 3

	13	14
70 Vinyl Acetate/30 PEG-8000 Graft Copolymer	100.0	90
Deodorized rosin		10
Glass Transition Temperature (Tg)	+5°C	+9°C
Viscosity at 177°C (350°F)	2750 mPas (cps)	975 mPas (cps)
Open Time	>15 sec	>15 sec
Adhesion to Tissue	F.T.	F.T.
Adhesion to polypropylene nonwoven	F.T.	Not tested
Spray Characteristics	Good	Good
"strike-through" on tissue at 16-20 g/m ²	good-no strike through	poor-strike through
21°C (70°F)/81% RH 3 days	none	
21°C (70°F)/81% RH 15 days	none	
peel (bonds aged 1 day)	100°F	
peel (bonds aged 1 week)	110°F	
shear (bonds aged 1 day)	110°F	
shear (bonds aged 1 week)	120°F	

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TABLE 3 (continued)

	13	14
Thermal Stability (72 hours/177°C) (350°F)		
Color	Amber	Amber
Skin	None	None
Gel	None	None
% viscosity	-17.3%	-25.1%

This example shows a hot melt suitable for nonwoven applications (13). It uses no tackifier and yet provides a good bond. The use of a tackifier (14) results in strike-through which is undesirable for this specific formulation. Hot melt formulation (13) demonstrates the useful property of delayed crystallization which diminishes "strike-through" and results in an increase in heat resistance over time. The incorporation of tackifying resin for many such systems (e.g., 14), results in a retardation of crystallization which is detrimental to bonding of multi-ply tissue.

EXAMPLE 4

In this example a graft polymer comprising 45 parts by weight vinyl acetate, 15 parts butyl acrylate, 25 parts polypropylene oxide/ethylene oxide (75H-90,000) and 15 parts polyethylene oxide (PEG-8000) was tested "neat" and formulated as an adhesive for water sensitive nonwoven applications.

Table 4

	15	16
45 Vinyl Acetate/ 15 Butyl Acrylate/ 25 U Conn 75 H/90,000 15 PEG-8000	100.0	70.0
Gum Rosin	---	30.0
Viscosity @ 177°C (350°F)	2,725 cps	Not Tested
Nonwoven Adhesion	0.39 N/cm (0.22 lb/in)	0.74 N/cm (0.42 lb/in)
Nonwoven/LDPE Adhesion	0.35 N/cm (0.20 lb/in)	0.91 N/cm (0.52 lb/in)
Water Sensitivity		
15 min/21°C (70°F)	No F.T.	No F.T.
24 hr/21°C (70°F)	No F.T.	No F.T.
15 min/38°C (100°F)	No F.T.	No F.T.
24 hr/38°C (100°F)	No F.T.	No F.T.
Heat Stability (72 hrs/177°C (350°F))		
Color	Dark Brown	Dark Brown
Skin	Partial	Full
Gel	None	None
% Viscosity Change	-61.1%	Not Tested

The neat polymer (15), exhibits acceptable adhesion to nonwoven and LDPE as well as water sensitivity. Incorporation of 30% resin (16) increases the adhesion values to both substrates without negatively affecting the water sensitivity and therefore such highly tackified formulations are particularly suitable for bonding of substrates such as in diaper construction. Although the color and skin formation are poorer than those reported in the prior examples, it is to be noted that no antioxidant was post added to either formulation. Addition of antioxidants such as are used by those skilled in the art will result in a reduction of skin formation.

Claims

1. A process for bonding nonwoven fabric substrates to similar or dissimilar substrates comprising the steps of applying to at least one substrate a molten hot melt adhesive composition, said adhesive having a Tg of -40 to +40°C

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comprising 50-100 weight % of a graft copolymer comprising (a) 40-85 weight percent of at least one vinyl monomer and (b) 15-60 weight percent of at least one polyalkylene oxide polymer having a molecular weight greater than 3000 and a polymerized ethylene oxide content of at least 50% by weight, and 0-50 weight % of a compatible tackifying resin; and bonding another substrate thereto.

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2. The process of Claim 1 wherein at least one substrate is a nonwoven and wherein the adhesives comprises 50-95 weight % of the graft copolymer and 5-50 weight % of a compatible tackifying resin.

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3. The process of Claim 1 wherein a superabsorbent particulate polymer is bonded to a nonwoven substrate and wherein the adhesive comprises 100% of the graft copolymer.

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4. The process of Claim 1 wherein the adhesive additionally comprises up to 25% by weight of a diluent selected from the group consisting of glycerol mono- and distearate, synthetic long chain linear polymeric alcohols, stearic acid, high acid number waxlike materials of mono- or dicarboxylic acids, acid wax derived from montan wax, stearyl alcohol, hydrogenated castor oil, ethoxylated alcohols, 12(OH) stearic acid and stearic-cetyl alcohol.

Patentansprüche

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1. Verfahren zum Verbinden von nicht-gewebten textilen Substraten mit ähnlichen oder verschiedenen Substraten, umfassend die Schritte: Auftragen einer geschmolzenen Schmelzkleberzusammensetzung auf mindestens ein Substrat, wobei der Kleber eine Tg von -40 bis +40°C aufweist und 50-100 Gewichts-% eines Pfropfcopolymers, umfassend (a) 40-85 Gewichtsprozent von mindestens einem Vinylmonomer und (b) 15-60 Gewichtsprozent von mindestens einem Polyalkylenoxidpolymer mit einem Molekulargewicht von mehr als 3000 und einem Gehalt von mindestens 50 Gewichts-% an polymerisiertem Ethylenoxid, sowie 0-50 Gewichts-% eines verträglichen klebrig-

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machenden Harzes umfaßt, und Binden eines anderen Substrats daran.

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2. Verfahren nach Anspruch 1, wobei mindestens ein Substrat ein nicht-gewebtes Textil ist und wobei die Kleber 50-95 Gewichts-% des Pfropfcopolymers und 5-50 Gewichts-% eines verträglichen klebrigmachenden Harzes umfaßt.

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3. Verfahren nach Anspruch 1, wobei ein superabsorbierendes, aus Partikeln bestehendes Polymer mit einem nicht-gewebten textilen Substrat verbunden wird und wobei der Kleber 100 % des Pfropfcopolymers umfaßt.

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4. Verfahren nach Anspruch 1, wobei der Kleber zusätzlich bis zu 25 Gewichts-% eines Verdünnungsmittels umfaßt, das aus der aus Glycerinmono- und Glycerindistearat, synthetischen langkettigen, linearen polymeren Alkoholen, Stearinsäure, wachsartigen Materialien mit hoher Säurezahl aus Mono- oder Dicarbonsäuren, von Montanwachs abgeleitete saure Wachse, Stearylalkohol, hydriertem Rizinusöl, ethoxylierten Alkoholen, 12(OH)-Stearinsäure und Stearyl-Cetylalkohol bestehenden Gruppe ausgewählt ist.

Revendications

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1. Procédé pour la liaison de substrats en étoffe non tissée à des substrats identiques ou différents, comprenant les étapes d'application à au moins un substrat d'une composition d'adhésif thermofusible fondue, ledit adhésif ayant une Tg de -40 à +40°C et comprenant 50 à 100 % d'un copolymère greffé comprenant (a) 40 à 85 pour cent en poids d'au moins un monomère vinylique et (b) 15 à 60 % en poids d'au moins un polymère d'oxyde d'alkylène ayant un poids moléculaire supérieur à 3000 et une teneur en oxyde d'éthylène polymérisé d'au moins 50 pour cent en poids, et 0 à 50 pour cent en poids d'une résine d'adhésivité compatible; et de liaison d'un autre substrat audit substrat.

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2. Procédé suivant la revendication 1, dans lequel au moins un substrat est un substrat non tissé et dans lequel l'adhésif comprend 50 à 95 pour cent en poids du copolymère greffé et 5 à 50 pour cent en poids d'une résine d'adhésivité compatible.

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3. Procédé suivant la revendication 1, dans lequel un polymère hyperabsorbant en particules est lié à un substrat non tissé et dans lequel l'adhésif comprend 100 pour cent du copolymère greffé.

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4. Procédé suivant la revendication 1, dans lequel l'adhésif comprend en outre jusqu'à 25 % en poids d'un diluant choisi dans le groupe consistant en mono- et distéarates de glycérol, alcools polymériques linéaires synthétiques à chaîne longue, acide stéarique, matières à haut indice d'acide analogue à des cires dérivées d'acides mono- ou dicarboxyliques, cire acide dérivée de la cire montan, alcool stéarylique, huile de ricin hydrogénée, alcools éthoxylés, acide 12(OH)-stéarique et alcool stéarique-cétylique.

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